08 February 2024
EMA/28356/2024
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): ferucarbotran

Procedure No. PSUSA/00001382/202306

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Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Resotran 540 mg/ml Injektionssuspension	SE/H/2120/001	7002837.00.00	B.E.IMAGING.GMBH	DE
Resotran 540 mg/ml injektionsvätska, suspension	SE/H/2120/01	61472	B.E.IMAGING.GMBH	SE